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## **Supplemental Material**

## Perinatal DDT Exposure Induces Hypertension and Cardiac Hypertrophy in Adult Mice

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Table S1. Water- and captopril- intake during the 1 week period of captopril administration.

	Male		Female	
	Vehicle	DDT	Vehicle	DDT
Mean water intake, water ml/kg	196 (8)	190 (8)	255 (6)	238 (6)*
body weight/day (SE)				
Mean captopril intake, captropril	112 (5)	108 (5)	125 (3)	117 (3)
mg/kg body weight, day (SE)				

<sup>\*</sup>p<0.05 DDT vs. vehicle controls within each sex

Table S2. Summary of ages and numbers of mice during experimental measurements.

ACEi, angiotensin converting enzyme inhibitor (captopril); BP, blood pressure measured by CODA.

Age	Sex	Measurement	Sample Size	Sample size
			(mice/litter)	(litter/treatment)
5	Male	BP	2	15 DDT,
				14 VEHICLE
5	Female	BP	2	15 DDT,
				14 VEHICLE
7	Male	BP, ACEi	2	7
7	Female	BP, ACEi	2	7
7	Male	telemetry	1	6 VEH+Water
				4 DDT+Water
				3 VEH+ACEi
				4 DDT+ACEi
7	Female	BP, renal transporter	1	6
		RNA		
8	Male	EchoMRI, renal	1	8
		pathology		
8	Female	EchoMRI, renal	1	8
		pathology		

Table S3. Decrease in systolic blood pressure (mmHg) by captopril ACE inhibition relative to water treated controls.

Method	Sex	Mean (SE) among	Mean (SE) among	Interaction
		Perinatal Vehicle	Perinatal DDT	p-value
Tail cuff	Female	22.6 (3.3)	33.9 (3.3)	0.02
Tail cuff	Male	21.5 (4.0)	32.7 (4.0)	0.05
Telemetry	Male	30.5 (0.5)	34.0 (0.3)	< 0.0001

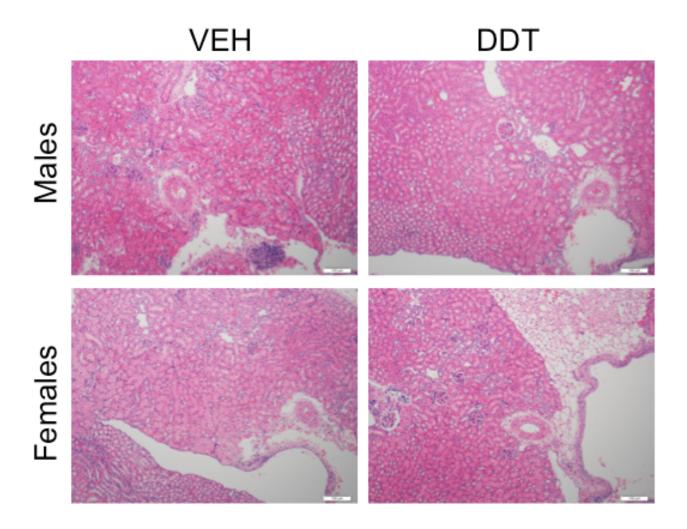


Figure S1. No renal pathology associated with perinatal DDT exposure. Kidney morphology as visualized by H&E stain looks normal in 8 month old mice. N=1 mouse/sex/litter and 8 litters/treatment.